AMERICAN HEALTH PACKAGING®
An AmerisourceBergen Company

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**URGENT PRODUCT RECALL**

April 8, 2019

Dear Valued Customer,

American Health Packaging, Inc. is initiating a voluntary drug recall to the **RETAIL LEVEL** for **AHP GlipiZIDE Extended-release Tablets 2.5 mg, 30 count Unit Dose Blisters, Carton NDC#: 68084-295-21, (Individual Dose NDC: 68084-295-11),** for the lot listed below:

| Product Description | AHP Lot No. | Expiration Date | Ship Dates of Product |
|---|---|---|---|
| **GlipiZIDE Extended-release Tablets 2.5 mg, 30 count Unit Dose Blisters**<br>**Carton NDC#: 68084-295-21**<br>**(Individual Dose NDC: 68084-295-11)** | 181288 | 5/31/2020 | 10/17/2018 to 12/31/2018 |

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| **REASON** | This recall is being initiated due to dissolution failure at time zero of the repackaged lot. |
| **HEALTH HAZARD EVALUATION** | Glipizide extended-release tablets are indicated as an adjunct to diet and exercise to improve glycemic control in adults with type 2 diabetes mellitus. |

**ACTIONS REQUIRED**

1. **Distributors/Pharmacies** - Immediately examine your inventory, quarantine and discontinue distribution of these lots.
2. **Distributors** - Complete the enclosed Business Reply Card even if you do not have any product on hand.
3. **Distributors** - Please pass this Recall Notice on **ONLY** to pharmacies that received these product lots.
4. **Pharmacies** - If you have units of the affected products/lots in inventory, please contact Inmar Rx Solutions at 877-475-5868 to receive a recall return packet. If you have not received this product you do not need to request a return packet.
5. **Distributors/Pharmacies** - Return recalled product to Inmar as instructed in recall/return packet.
6. **Pharmacies** - You do not need to contact any patients.
7. Carry out a physical count and record this data on the Business Reply Card and the Packing slip, which are included with this letter.
8. Return the recalled product and the Packing Slip using the prepaid Fedex shipping labels to:

   **Inmar Rx Solutions Inc**
   **6101 North 64th Street**
   **Milwaukee, WI 53218**

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|---|---|
| **OTHER** | This Recall extends to the Retail Level only. No other lots, packages, or formulations are being recalled.<br>Please reorder stock immediately.<br>For questions about the recall process, call Inmar Rx Solutions at (877) 475-5868<br>This recall is being conducted with the knowledge of the Food and Drug Administration. We appreciate your immediate attention and cooperation and sincerely regret any inconvenience caused by this action. |

**To receive credit, the reply form and recalled product must be returned to GENCO by July 10th, 2019.**

Thank you for your support in complying with the requests in this letter. We apologize for the inconvenience that this incident may cause you or your customers.

Sincerely,

Becky A Mahon

Becky A Mahon
Regulatory Specialist
American Health Packaging